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AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Original) A dosage form of dalbavancin for parenteral use comprising:
- a sterile, stable, particle-free dalbavancin powder suitable for reconstitution with a pharmaceutically acceptable vehicle; and

a stabilizer,

wherein the dosage form is at a pH of about 3-5.

- 2. (Original) The dosage form of claim 1, wherein the stabilizer comprises sugar.
- 3. (Original) The dosage form of claim 2, wherein the sugar is selected from the group consisting of mannitol, lactose, sucrose, sorbitol, glycerol, cellulose, trehalose, maltose, dextrose, and combinations thereof.
 - 4. (Original) The dosage form of claim 1, wherein the stabilizer is mannitol.
- 5. (Original) The dosage form of claim 4, wherein the weight ratio of mannitol: dalbavancin is 1:2.
 - 6. (Original) The dosage form of claim 1, wherein the stabilizer is lactose.

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- 7. (Original) The dosage form of claim 1, wherein the stabilizer is a mixture of mannitol and lactose.
- 8. (Original) The dosage form of claim 7, wherein the weight ratio of mannitol:lactose:dalbavancin is 1:1:4.
 - 9. (Original) The dosage form of claim 8, wherein the pH is 4.5.
 - 10. (Original) The dosage form of claim 1, wherein the pH is 3.5.
 - 11. (Original) The dosage form of claim 1, wherein the pH is 4.5.
 - 12-24. (Canceled)
- 25. (Original) A dosage form of dalbavancin for parenteral use comprising:

 a sterile, stable, particle-free dalbavancin powder suitable for reconstitution with a pharmaceutically acceptable vehicle and mannitol at a pH of about 3-5.
- 26. (Original) The dosage form of claim 25, wherein the pharmaceutical composition further comprises lactose.
 - 27. (Original) The dosage form of claim 25, wherein the pH is about 3.5.

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28-32. (Canceled)

- 33. (Original) A pharmaceutical composition comprising:dalbavancin; anda stabilizer, wherein the stabilizer comprises mannitol and lactose.
- 34. (Original) The pharmaceutical composition of claim 33, wherein the weight ratio of mannitol:lactose:dalbavancin is 1:1:4.
- 35. (Original) The pharmaceutical composition of claim 33, wherein the pharmaceutical composition has a pH of about 3 to 5.
- 36. (Original) The pharmaceutical composition of claim 33, wherein the pharmaceutical composition has a pH of about 4.5.

37-66. (Canceled)

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